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(54) Enhanced yeast transcription employing hybrid promoter region constructs.

(57) Yeast promoters of glycolytic enzymes are modified by isolating a fragment encompassing the RNA polymerase binding site and joining to the 5' end of this fragment a DNA sequence providing for enhanced inducible or constitutive transcription of a structural gene. Constructs are prepared for efficient expression of foreign genes in yeast.

Yeast strains 2150-2-3(pCI/1GAPSOD) and AB110(pCI/1GAPAT19), producing human α -antitrypsin and superoxide dismutase, were deposited at the A.T.C.C. on May 9, 1984 and given Accession Nos. 20708 and 20709, respectively; and 2150-2-3(GAP5), 2150-2-3(Pyk5) and 2150-2-3(PHO5GAP1), expressing Hepatitis B surface antigen, were deposited at the A.T.C.C. on May 9, 1984 and given Accession Nos. 20705, 20706 and 20707, respectively.

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**ENHANCED YEAST TRANSCRIPTION EMPLOYING HYBRID
PROMOTER REGION CONSTRUCTS**

BACKGROUND OF THE INVENTION

5 Field of the Invention

With the advent of hybrid DNA technology, production of mammalian proteins in microorganisms became a reality. For the first time, mammalian proteins could be synthesized in a unicellular microorganism by
10 introduction of a gene encoding for the mammalian protein under the transcriptional and translational control of regulatory sequences recognized by the microorganism host. The introduction of these foreign constructions into a microorganism host resulted in competition be-
15 tween the regulatory signals of the construct and the regulatory signal endogenous to the host for the host systems involved with expression. The structural gene of interest is usually directed to a product which is nonproductive and may be injurious to the host. There-
20 fore, host cells which can turn off the foreign gene can effectively dominate modified host cells.

Substantial progress has been made in isolating sequences concerned with transcriptional and translational regulation for protein expression. But frequently flanking sequences, as well as distant sequences,
25 may also affect the efficiency and regulation of the expression of the protein. Therefore, as one manipulates these various sequences, removing them from their native environment, and joining them to unnatural sequences,
30 that is sequences other than the wild-type sequence, one can never be certain as to the result.

In order to enhance the economies of producing proteins in microorganisms, there have been substantial efforts directed to improving the efficiency of tran-
35 scription and translation, maximizing the prop rti n f

total protein directed to production of the desired product, enhancing the viability of the modified host, as well as improving the efficiency with which the modified host may be obtained.

5 Description of the Prior Art

Guarente et al., Proc. Natl. Acad. Sci. USA (1982) 79:7410-7414, describes a hybrid promoter region employing the GAL4 regulatory region. Guarente and Ptashne, ibid. (1981) 78:2199-2203, report the presence
10 of two domains in a yeast promoter, with a region upstream from the TATA sequence providing an activation site. Kramer et al., ibid. (1984) 81:367-370, describe the regulated expression of a human interferon gene in yeast employing the yeast acid phosphatase promoter,
15 where expression is induced by phosphate or a temperature shift. Tekamp-Olson et al., Cold Spring Harbor Meeting, Molecular Biology of Yeast, 1983, describe the absence of deleterious effects on yeast viability when employing "short" promoters, as distinct from the presence
20 of such effects, when employing an extended promoter region or "long" promoters.

SUMMARY OF THE INVENTION

Novel hybrid promoter regions are provided for use in conjunction with constructs having a structural gene under the transcriptional control of the
25 hybrid promoter region and a terminator region. The hybrid promoter comprises a first segment providing transcriptional enhancement, either constitutive or regulated, and a second segment defining an RNA polymerase binding site and transcriptional initiation site.
30 The hybrid promoter regions lack the deleterious effects associated with a wild-type promoter region in recombinant constructs, which results in reduced transformation efficiencies and lower yeast viability.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a diagrammatic view of plasmids GAP1-6;

Figur 2 is a diagrammatic view f plasmids Pyk1-6; and

Figure 3 indicates the DNA linker sequence and a flow diagram showing its use in a construct for hSOD.

DESCRIPTION OF THE SPECIFIC EMBODIMENTS

10 Novel DNA fragments are provided as well as constructions employing the fragments for enhanced expression of structural genes in a yeast host. The constructs employing a hybrid promoter region provide for enhanced efficiencies of transformation and greatly
15 improved viability of the yeast host as contrasted with those employing a wild-type yeast promoter. Concomitant with the improved viability is increased expression of a structural gene, in comparison with the truncated promoter region, and, therefore, greatly enhanced over-
20 all yields of expression products.

For the purposes of the subject invention, the "promoter region" is divided into two domains: (1) the structural gene proximal region, which includes the transcription initiation site, the "TATA" sequence
25 capping sequence, as appropriate, and an RNA polymerase binding sequence, which sequence intends a sequence which includes nucleotides upstream from the initiation site for directing the initiation of synthesis of the messenger RNA; and (2) a distal region which provides
30 for regulated or constitutive expression, with enhanced expression as compared to the first domain linked to non-functional yeast DNA.

The hybrid promoters of the subject invention employ the RNA polymerase binding region of a yeast
35 glycolytic enzyme promoter and a r gi n upstr am from said polymerase binding region, which is different from

th wild-type upstream region of the RNA polymerase binding region and provides for enhanced efficiencies of transcription. This distal region will be derived from either a sequence, usually a yeast sequence, involved in regulation of transcription, or a prokaryotic sequence which provides for enhanced constitutive expression of the desired gene.

Conveniently, cassettes or constructs can be prepared which provide for one or more restriction sites intermediate the promoter region and a related terminator region where the structural gene may be inserted, so as to be under the transcriptional control of the hybrid promoter region. By having one or more restriction sites, one can provide for ease of insertion of the structural gene intermediate the transcription initiation and termination regions. The cassettes which can be prepared comprising the transcriptional initiation and termination region, having appropriate restriction sites for structural gene insertion can be cloned in prokaryotic vectors, so that after insertion of the structural gene, the resulting cassette, including the structural gene, may be cloned, isolated and purified, before introduction into a yeast vector.

The cassette, will for the most part, have the following formula:

-(P.R.(2)-P.R.(1))-R.S.-T.R.-

wherein:

P.R.(1) is the promoter region proximal to the structural gene and having the transcription initiation site, the RNA polymerase binding site, and including the TATA box, the CAAT sequence, as well as translational regulatory signals, e.g., capping sequence, as appropriate;

P.R.(2) is the promoter region joined to the 5'-end of P.R.(1) associated with enhancing the

efficiency of transcription of the RNA polymerase binding region;

R.S. is a sequence having one or more restriction recognition sites, preferably at least two restriction recognition sites, where the sites may result upon restriction into blunt ends or overhangs;

T.R. intends the termination region, which will include the terminator, which may be a stem and loop structure, and which may be associated with one or more stop codons, a polyadenylation signal sequence, if any, as well as any other transcriptional and translational termination sequences.

P.R.(1) will generally be at least about 150bp, more usually at least about 200bp, usually not more than about 600bp, more usually not more than about 500bp, generally not more than about 450bp and preferably less than about 400bp; the sequence will extend in the downstream direction of transcription to about bp +3, more usually bp -1 and may extend only to bp -20, more usually to bp -10 (the numbering intends that +1 is the first bp of the initiation codon with which the promoter region is associated in the wild-type host, while -1 is the immediately upstream bp and the integers increase in the direction of transcription;

P.R.(1) will be derived from a strong yeast promoter, normally a glycolytic enzyme promoter, such as glyceraldehyde-3-phosphate dehydrogenase, pyruvate kinase, alcohol dehydrogenase, phosphoglucosomerase, triose phosphate isomerase, phosphofructokinase, etc.;

P.R.(2) will provide for an enhancing function of transcription, which enhancing function may provide for constitutive or regulated transcription; regulators will be derived from regions associated with regulation of yeast genes, other than the natural or wild-type gene associated with the first domain in the wild-type or natural host, such as UDP-galactose epimerase (GAL10), galactokinase (GAL1), acid phosphatase (PHO5), alcohol

dehydrogenas I and II (ADH I and II), etc. For y ast regulatory sequences, the domain will usually be at least about 100bp, mor usually at least ab ut 200bp, for convenience generally not exceeding about 3kbp, usually not exceeding about 1kbp, desirably not exce d-
5 ing about 600bp. The regulatory region will generally begin at least about 200bp from the initiation cod n, usually at least about 300bp and may begin at 400bp or farther upstream from the initiation codon.

10 Regulation can be as a result of a change in the chemical or physical environment of the host, such as a change in carbon source, e.g., glucose to galactose or vice versa; a change in concentration of a nutrient, e.g., an inorganic nutrient such as a phosphate; or a
15 change in temperature, e.g., 25°C to 35°C. Constitutive transcription can be achieved employing prokaryotic sequences of at least about 500bp, usually 1kbp or more, for convenience, generally not exceeding about 5kbp; conveniently, the prokaryotic sequence can be obtain d
20 from the vector in which the cassette is cloned, illustrative vectors including pBR322, lambda, Charon 4A, pACYC184, pUC5, etc.

R.S. will generally be at least 4bp, more usually at least 6bp, and may be 100bp or more, more
25 usually being not more than about 60bp and may include one or more, usually not more than about 10 restriction sites, where such restriction sites may be illustrated by EcoRI, BamHI, Sall, HindIII, AluI, AvaI, TaqI, HpaI, etc., having at least one unique restriction site for
30 the construct sequences.

T.R. is the termination region which will include the necessary transcriptional and translational signals for termination, such as the polyadenylation site, etc.;

35 T.R. will generally be at least about 100bp, m r usually at least 150bp, and g nerally less than

5

Where a cassette is cloned in a bacterial vector, the construction will have the following formula:

10

wherein all the symbols have been defined previously,
except for:

15

20

30

such as cup^+ genotyp ; r sistanc to antibi tics, such as amp^r , tc^r , cam^r , str^r , tur^r genotyp , tc.;

b is 0 or 1, intending that the constructi n is either linear or circular, usually circular.

5 The above construct can be used for insertion of a wide variety of structural genes, both prokaryotic and eukaryotic, both naturally occurring and synth tic, where the genes may include signal leaders for secretion, and the like. The genes may express enzymes, hormones,
10 proteins from pathogens for vaccines, structural prot ins, lymphokines, membrane surface proteins, immunoglobulins, blood proteins, or the like. The particular structural gene which is inserted is not critical to this invention and any polypeptide or protein of interest may be pre-
15 pared employing the constructions of the subject inven- tion. The structural genes will usually be foreign to the yeast host, where foreign intends different fr m wild-type yeast structural genes and from a source that does not normally exchange genetic information with
20 yeast.

Usually, the structural gene will be at least about 36bp, and not more than about 20kbp, usually not more than about 3000bp, usually not more than about 1500bp. Included in the structural gene may be non-
25 coding flanking regions, the 5'-flanking region normally being quite short, usually less than about 30bp, whil the 3'-flanking region may be extended, usually not exceeding about 500bp. Thus, the structural gene frag-
30 ment will usually include the translational stop c dons for proper termination of amino acid chain extension.

When the structural gene has been inserted into the cassette which is joined to a yeast replication system, normally including one or more markers recog-
nized by yeast, the resulting construct will have the
35 following f rmula:

-(P.R.(2)-P.R.(1))-gene-T.R.-Rep(Y)-(M(Y))_a-

where in all of the symbols have been defined previously except for:

5 gene, which intends the structural gene, having its initiation codon and stop codons as appropriate; and

 Y, which intends that the symbol is related to yeast.

10 Convenient yeast replication systems include the 2µm plasmid replication system, combination of CEN3 and ARS1 or ARS3, or the like. The replication systems may be high or low copy number, depending on the effect of the construct on the viability of the host. While
15 the indicated replication systems are those which have found common employment, any replication system useful in yeast may be employed which provides for efficient replication and maintenance. Often the structural gene will be inserted into an appropriate shuttle vector
20 capable of replication and selection in either a yeast or bacterial host, where the resulting construction will have the following formula:

-(P.R.(2)-P.R.(1))-gene-T.R.-Rep(Y)-(M(Y))_a-Rep(B)-(M(B))_a-

25 where all symbols have been defined previously. Also, it is, of course, understood that the cassette without an inserted structural gene but containing the restriction enzyme recognition sequence, R.S., may be propagated in yeast or contained within a shuttle vector, where
30 the construction will have the following respective formulae:

-(P.R.(2)-P.R.(1))-R.S.-T.R.-Rep(Y)-(M(Y))_a-

-(P.R.(2)-P.R.(1))- R.S.-T.R.-Rep(Y)-(M(Y))_a-Rep(B)-(M(B))_a-

where all symbols have been defined previously.

The various fragments which form the cassette
 5 and final constructions may be joined together in accordance with conventional ways. In many cases, genes have been isolated and restriction mapped, as well as sequenced. To that extent, one can select the sequence of interest by restriction of the gene, employing further
 10 manipulation as necessary such as resection with Bal31, in vitro mutagenesis, primer repair, or the like, to provide a fragment of a desired size, including the desired sequence, and having the appropriate termini. Linkers and adapters can be used for joining sequences,
 15 as well as replacing lost sequences, where the restriction site is internal to the region of interest. The various fragments which are isolated, may be purified by electrophoresis, electroeluted, ligated to other sequences, cloned, reisolated and further manipulated.

20 The use of regulatory sequences for controlling transcription of the structural gene of interest allows for growing the host cells to high density with no or low levels of expression of the structural gene, and then inducing expression by changing the environmental
 25 conditions, e.g., nutrient, temperature, etc.

For example, with the GAL4 regulatory region, the yeast cells could be grown in rich media with a glycerol-lactic acid combination to high density, e.g., mid or late log phase, followed by switching the carbon
 30 source to galactose. For PHO5 regulation one could grow the cells at high phosphate, about 1 to 10mM, and then decrease the phosphate concentration to about 0.1 to 0.5mM. For temperature sensitivity, one could grow the cells at 25° to 37°C and then change the temperature
 35 as appropriate by about 5° to 20°C. The host cells would have the regulatory system associated with the regulatory region employed.

Various techniques will be exemplified in the Experimental section of this application, which techniques can be used as paradigmatic for constructions employing fragments from sources other than those exemplified. Of particular interest, as evidenced by the Experimental section, will be the use of the glyceraldehyde-3-phosphate dehydrogenase promoter region for the RNA polymerase binding site in conjunction with regulator sequences, such as those associated with GAL4, PHO5, ADHII or the like. In referring to the GAL4 regulatory region or associated sequence, the region intends the sequence associated with regulation of other galactose metabolism genes, e.g., GAL1 and GAL10, which are under the regulatory control of such sequence in conjunction with the expression product of the GAL4 gene. The PHO5 and ADHII sequences refer to regions associated with the PHO5 and ADHII genes which provide for transcriptional regulation of the PHO5 and ADHII gene, respectively.

The following examples are offered by way of illustration and not by way of limitation.

EXPERIMENTAL

All DNA manipulations were done according to standard procedures. See Molecular Cloning, T. Maniatis et al., Cold Spring Harbor Lab., 1982. Enzymes used in cloning were utilized as per the manufacturer's specifications. Enzymes were obtained either from New England Biolabs or Bethesda Research Laboratories. Procedures with these enzymes employed the supplier's directions.

Yeast were transformed and grown using a variety of media including selective medium (yeast nitrogen base without leucine); YEPD medium, containing 1% (w/v) yeast extract, 2% (w/v) peptone and 2% (w/v) glucose, and others as appropriate and/or detailed below. In the case of plating medium contained 2% (w/v) agar and for transformation 3% top agar. Hepatitis B surface antigen

was determined after lysis of yeast by glass bead agitation and clarification by centrifugation, using the AusriaII assay (Abbott Laboratories). Protein is determined by the Coomassie dye binding method.

5 Construction of GAL regulator containing plasmids.

Plasmid pLGSD5 is prepared as described in Guarente et al., (1982) supra. The plasmid was manipulated as follows: After restriction with XhoI, the overhangs were filled in with the Klenow fragment of
10 DNA polymerase I ("Klenow fragment"), ligated with EcRI linkers (GGAATTCC) and then completely digested with EcoRI and Sau3A to provide a 370bp fragment which was isolated by gel electrophoresis and included the intergenic sequence between GAL1 and GAL10 genes of yeast,
15 and provides for the GAL4 regulation sequence of the GAL1 and GAL10 genes.

This fragment was inserted into pBR322 which had been completely digested with EcoRI and BamHI, followed by treatment with alkaline phosphatase to prevent
20 oligomerization. The resulting plasmid pBRGAL4 was treated in two different ways.

In the first procedure pBRGAL4 was completely digested with Sau3A, the overhangs filled in with the Klenow fragment, and the resulting blunt-ended fragment
25 ligated with SalI linkers (CGTCGACG), followed by digestion with SalI and XhoI. The resulting 370bp fragment was isolated by gel electrophoresis. This fragment has the original 370bp yeast GAL4 regulator sequence with XhoI and SalI termini.

30 The second fragment was obtained by complete digestion of pBRGAL4 with XhoI and SalI to provide a XhoI-SalI fragment which included the 370bp yeast GAL4 regulator sequence as well as about 280bp of pBR322, the GAL4 sequence extending from Sau3A to SalI.

The two fragments were then cloned in the plasmid p1 t5. p1 t5 was prepared by inserting the 40bp polylinker fragment of the following sequence



- 10 into pBR322 as an EcoRI-PvuII substitution followed by insertion of the trp-lac promoter (Russell and Bennett, Gene (1982) 20:231-245) into the PvuII site with transcription oriented toward the polylinker sequence.
- 15 plot5 was completely digested with SalI, followed by treatment with alkaline phosphatase and the 370bp and 650bp fragments independently inserted into plot5 to provide plasmids plot5GAL4/370 and plot5GAL4/650, respectively. Each of the plasmids was then completely digested with BamHI and SalI to reproduce the individual
- 20 fragments extended by 6bp of the polylinker fragment. These fragments were then ligated into pC1/1, which had been completely digested with BamHI and SalI followed by treatment with alkaline phosphatase to prevent recircularization. Plasmid pC1/1 is a derivative of
- 25 pJDB219 (Beggs, Nature (1978) 275:104) in which the region corresponding to bacterial plasmid pMB9 in pJDB219 has been replaced by pBR322 in pC1/1. The resulting plasmids were designated pC1/1GAL4/370 and pC1/1GAL4/650, respectively. The BamHI-SalI fragment
- 30 is located in the pBR322 portion of the vector pC1/1.

The next construction develops a hybrid promoter for expression of the Hepatitis B surface antigen (HBsAg or sAg), employing the RNA polymerase binding region of GAPDH. The previously prepared plasmid

pHBS56/16-3, a yeast shuttle vector containing the alcohol dehydrogenase 1 (ADH1) promoter, the HBsAg gene and ADH terminator as a SphI fragment, was digested with SphI and the ends modified with Bam linkers. The
5 Bam linkers have the sequence CGGATCCG.

pHBS56/16-3 was prepared as follows: A TagI-HpaI fragment obtained from the HBsAg coding region which included 26bp of the pre-sAg region, 681bp of the sAg region and 128bp of the 3'-untranslated region, was
10 linked with EcoRI linkers and cloned at the EcoRI site in pBR322. The EcoRI linkers have the sequence GGAATTCC. The plasmid pHBS5 was thus obtained.

After digesting pHBS5 with EcoRI, the digest was resected with Bal31 and religated with EcoRI linkers
15 (GGAATTCC). After digestion with EcoRI the material of about 800bp was isolated from a polyacrylamide gel. This isolate was then recloned into pBR322 which had been digested with EcoRI and treated with alkaline phosphatase. Where the resection was to the sequence CATGG,
20 which included the methionine codon, the EcoRI linker created an NcoI site. The plasmids were screened for the presence of an NcoI site and one of the plasmids chosen for further manipulation. This plasmid, designated pHBS5-3, was restricted with EcoRI, the EcoRI
25 fragment made blunt-ended with Klenow fragment and dNTPs, and the blunt-ended fragment was then restricted with XbaI to provide an about 100bp fragment having an XbaI overhang and blunt end at the former EcoRI site.

pHBS5 was then digested with ClaI, made blunt-ended with the Klenow fragment and dNTPs, digested with XbaI, followed by alkaline phosphatase treatment. The
30 100bp fragment was then inserted into the vector to provide the plasmid pHBS6. Upon sequencing of the blunt-ended ligation site, it was found that an adenosine had been lost, so as to lose the EcoRI site, where
35 the sequence was now ATCGATTCCCATGG. The ClaI and NcoI

sites were retained. The loss of the A resulted in pHBS6 having a single EcRI site.

pHBS5-3 was digested with EcoRI and the resulting EcoRI fragment having the sAg fragment isolated by gel electrophoresis and inserted into the vector pHBS16 (Valenzuela *et al.*, Nature (1982) 298:347-350). This plasmid has the ADH1 promoter and the sAg gene in an EcoRI fragment in a plasmid containing the 2 μ m origin, a TrpI gene and pBR322. The plasmid was digested with EcoRI, treated with alkaline phosphatase to prevent recircularization, and the EcoRI fragment from pHBS5-3 inserted to provide pHBS16-3, where the sAg gene isolated as a TagI-HpaI fragment had been modified by Bal31 resection. The plasmid pHBS16-3 was digested with SphI and XbaI to provide a fragment which had the ADH promoter at the Sph terminus and the 5'-end of the sAg gene.

pHBS56 was then digested with SphI. pHBS56 was prepared from pC1/1 by digestion with SphI, which deletes a portion of the plasmid spanning the 2 μ m-pBR322 joint. The active portion of the ADH1 promoter region is contained within the SphI-HindIII fragment of approximately 300bp (Bennetzen *et al.*, J. Biol. Chem. (1982) 257:301). The SphI site in the ADH promoter begins at position -413 and the yeast terminator sequence is contained within a HindIII-SphI fragment of about 330bp. In each case the SphI site is distal to the coding region. A 1500bp ADH1 promoter fragment terminating at position -9 (Hitzeman *et al.*, Nature (1981) 293:717) and an approximately 450bp terminator unit from nucleotides 913 to 1368 in the ADH gene nucleotide sequence were joined at a HindIII site between the fragments and cloned into the BamHI site of the vector YEpl3 (Broach and Hicks, Gene (1979) 8:121) to provide pADH5.

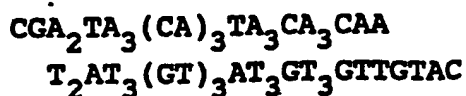
The HBsAg-DNA segment of pHBS5 was excised by EcoRI digestion, blunt-ended with the Klenow fragment and joined at both ends with HindIII linkers, CAAGCTTG. After digestion with HindIII, the HBsAg fragment was

ins rted into th HindIII site of the plasmid pADH5 which had b en .digested at the HindIII site intermediate the ADH1 promoter and terminator s quenc . A plasmid with the HBsAg gene in the correct orientation as determined by restriction analysis was designated pHBS22. The cassette was included between two SphI restricti n sites. pHBS22 was digested with SphI to obtain a fragment of about 1500bp and inserted into SphI digested pC1/1 to provide pHBS56 which was cloned in E. coli HB101.

pHBS56 was digested with SphI and XbaI to provide a 1.1kb fragment having the ADH terminator r - gion and the 3'-portion of the sAg gene with the SphI site proximal to the terminator region. The 1.1kb SphI-XbaI fragment was joined to the SphI-XbaI fragment from pHBS16-3, which resulted in providing the compl te sAg gene in the correct orientation between the ADH promoter and terminator. This SphI-SphI fragment was then ligated to SphI digested pHBS56, replacing th cassette of pHBS56 to provide the plasmid pHBS56/16-3 with the resected sAg coding region fragment. The cassette was then excised from pHBS56/16-3 by digestion with SphI, followed by chewing back the overhangs with the Klenow fragment in the presence of dNTPs, then ligated with BamHI linkers, followed by digestion with BamHI to provide a 1.6kb fragment which was isolated by gel electrophoresis. The fragment included the ADH promoter region, the sAg gene and ADH terminator region, as described above. This fragment was inserted into the BamHI site of pBR322 to provide pPGT16-3 which was digested with BamHI and XbaI and the resulting 1.1kb fragment gel isolated, where the XbaI-BamHI fragment had the 3' portion of the sAg gene and the ADH terminator region.

pHBS6 was digested with XbaI and Nc I and the 94bp fragment gel isolat d to provide th 5'-p rtion of

th sAg gene. A synthetic adapter was prepared of the formula



- 5 having TaqI and NcoI termini and providing the -25 to -1 nucleotides of the GAPDH (GAP49) promoter and the initiation codon of the sAg gene. This synthetic fragment, the NcoI-XbaI fragment, and the XbaI-BamHI fragment were ligated simultaneously, followed by digestion
10 with TaqI and BamHI. The resulting fragment was then substituted into pBR322 linearized with ClaI and BamHI, followed by treatment with alkaline phosphatase. The resulting plasmid, which contains the -1 to -25bp of the GAPDH promoter region, the sAg gene, and the ADH
15 terminator, where the NcoI restriction site is lost was called pHBS6LGAPsAgtADH.

- pGAP1, a plasmid prepared by insertion of a HindIII fragment containing the GAPDH gene GAP49 (Holland and Holland, J. Biol. Chem. (1979) 254:5466-5474) in-
20 serted in the HindIII site of pBR322, was digested with HinfI and a 500bp promoter containing fragment isolated. The fragment was resected with Bal31 to remove about 50 or 90bp, followed by ligation with HindIII linkers and digestion with HindIII. pBR322 was digested with
25 HindIII, followed by treatment with alkaline phosphatase and the about 450 or 410bp fragment inserted to provide pGAP128 and pGAP396, respectively.

- pGAP128 was digested with HindIII, the fragment made blunt-ended with the Klenow fragment and dNTPs and
30 the resulting 450bp fragment isolated by gel electrophoresis. This fragment was inserted into SmaI digested plot5, which had been treated with alkaline phosphatase, to provide plasmid plot5pGAP128, which contained about -400 to +27bp of the GAPDH promoter and coding region.
35 Plasmid plot5pGAP396 was prepared from pGAP396 in an identical manner and thus differs from plasmid

pl t5pGAP128 in having about 15-30 fewer bp at each terminus of the GAPDH promoter region (about -385 to -3).

Plasmids GAP1-GAP4 were then prepared in the following manner. Plasmid plot5pGAP128 was digested with TaqI and BamHI to provide an about 390bp TaqI-BamHI fragment which included the -26 to about -400bp of the GAPDH promoter region and a portion of the HindIII and plot5 polylinker. pHBS6LGAPsAgtADH plasmid was also digested with TaqI and BamHI and a 1.1kb TaqI-BamHI fragment containing the 3'-terminus of the GAPDH promoter region, the sAg gene and the ADH terminator region was gel isolated and ligated to the other TaqI-BamHI fragment to provide a BamHI-BamHI fragment which included approximately 400bp of the GAPDH promoter region, the sAg gene in proper orientation for transcriptional regulation by the GAPDH promoter, followed by the ADH terminator region. This fragment was ligated into pBR322 which had been digested with BamHI and treated with alkaline phosphatase to provide plasmid pPGT80. This BamHI cassette could now be isolated and inserted into plasmid pC1/1, at the BamHI site in the pBR322 portion of pC1/1, where in plasmid GAP1 the ADH terminator region is proximal to the amp^r gene with the pBR322 portion divided into an approximately 4kb sequence including the amp^r gene and a 375bp region separating the cassette from the 2 μ m sequences. In GAP2, the promoter is adjacent to the long pBR322 sequence with transcription in the same direction as the amp^r gene. The same cassette was inserted into BamHI-digested pC1/1GAL4/650 to obtain plasmids GAP3 and GAP4, where GAP3 has the GAPDH promoter distal from the GAL4 regulator region and the long pBR322 sequence and GAP4 has the GAPDH promoter adjacent to the GAL4 regulator region, which is adjacent to the long pBR322 sequence.

Plasmids GAP5 and GAP6 were isolated as follows. Plasmid plot5pGAP396 was digested with SalI and TaqI

and a fragment containing 9bp of the plot5 polylinker sequence and the GAPDH promoter sequence extending from about -385 to -26bp was isolated. An approximately 130bp TagI-XbaI fragment including -25 to -1bp of the GAPDH promoter and +1 to +93bp of the sAg gene was obtained from pHBS6LGAPsAgtADH. A 1.1kb XbaI-SalI fragment containing the 3'-portion of the sAg gene and the ADH terminator as well as 6bp of plot5 polylinker sequence was obtained from plasmid plot5sAgtADH (described below--Pyruvate Kinase Promoter). These three fragments were ligated, digested with SalI and then cloned into SalI-digested pC1/1GAL4/370. GAP5 has the GAPDH promoter region adjacent to the GAL4 regulator region, which is proximal to the short pBR322 sequence, and GAP6 has the GAPDH promoter region distal from the GAL4 regulator region and proximal to the long pBR322 sequence (see Fig. 1).

Pyruvate kinase promoter.

Plasmid pHBS6Pyk containing the sAg gene under the transcriptional regulatory control of the Pyk promoter was obtained by cloning a 4.4kb insert of yeast genomic DNA in pBR322 containing the Pyk gene and 911 nucleotides of 5'-untranslated region, and digestion of this plasmid pPyk9.1.1 with XbaI. After making the ends blunt-ended, the linear fragment was digested with BamHI providing a 912bp BamHI-blunt fragment containing the Pyk promoter and 8 bases from the Pyk coding region. This fragment was inserted into the plasmid pHBS6, which had been digested with NcoI, blunt-ended and digested with BamHI. The plasmid pHBS6Pyk was totally digested with EcoRI, to obtain a fragment including the sAg gene and a portion of the Pyk promoter region. The fragment was made blunt-ended with the Klenow fragment and dNTPs, followed by ligation to BamHI linkers, digested with XbaI, which is internal to the sAg gene, the XbaI terminus made blunt-ended with the Klenow

fragment and dNTPs, followed by digestion with BamHI, to provide a 580bp BamHI-blunt-ended (XbaI) fragment. The plasmid plot5 was digested with EcoRI, made blunt-ended, digested with BamHI and treated with alkaline phosphatase and the two fragments joined to provide

5 plasmids plot5PyksAg51 and plot5PyksAg.57. The two differ in that the BamHI site of the latter was not regenerated during cloning, possibly as a consequence of minimal nuclease contamination (digestion).

10 plot5 was treated as previously described (EcoRI digestion, blunt-ended, BamHI digestion and treatment with alkaline phosphatase) and joined to a 1.1kb fragment obtained by digestion of pPGT16-3 with XbaI, followed by blunt ending, followed by digestion with

15 BamHI and gel isolation. This fragment was introduced into plot5 to provide the plasmid plot5sAgtADH. Again the BamHI site in this plasmid was not regenerated, presumably due to digestion by contaminating nucleases.

Plasmids Pyk1 and Pyk2 were prepared as follows. Plasmid plot5PyksAg51 was digested with BamHI, then with XbaI, and an approximately 580bp fragment containing about 480bp of Pyk promoter and 93bp of the 5'-end of the sAg gene was gel isolated. A 1.1kb XbaI-SalI fragment containing the 3'-portion of the sAg gene, the ADH terminator and about 6bp of the plot5 polylinker

25 was isolated from plot5AgtADH. These two fragments were ligated, digested with BamHI and SalI and then cloned into plasmid pC1/1, which had been cleaved with BamHI and SalI and treated with alkaline phosphatase, to yield plasmid Pyk1. Plasmid Pyk2 was prepared similarly but the 580bp SalI-XbaI, Pyk promoter/HBsAg gene 5'-end fusion fragment was isolated from plot5PyksAg.57 and included about 6bp of plot5 polylinker sequence

30 upstream from the promoter region. Also the 1.1kb XbaI-BamHI fragment containing the 3'-part of the HBsAg gene and the ADH terminator was derived from plasmid pPGT16-3.

Plasmids Pyk3-Pyk6 were prepared as follows.

Plasmid plot5PyksAg51 was digested with BamHI, then with XbaI and the about 580bp fragment containing the Pyk promoter and the 5'-part of the HBsAg gene isolated as above. The 1.1kb BamHI-XbaI fragment, containing the 3'-portion of the HBsAg gene and ADH terminator, was recovered from pPGT16-3, also as above, and the two fragments ligated, digested with BamHI and inserted with different orientations into the BamHI site of pCl/1GAL4/650 (Pyk3, Pyk4). Plasmids Pyk5 and Pyk6 were prepared similarly except that the SalI-XbaI fragment containing the Pyk promoter and 5'-end of the sAg gene was isolated from plot5PyksAg.57 and the XbaI-SalI sAg gene 3'-portion/ADH terminator fusion fragment was derived from plot5sAgADH and thus both fragments included approximately 6bp of plot5 polylinker sequence. The cassette so formed was then cloned into the SalI site of pCl/1GAL4/370 in opposite orientations.

The six plasmids designated Pyk1-6 (see Fig. 2) are distinguished by Pyk1 having the promoter region proximal to the short pBR322 sequence; Pyk2 having the promoter region proximal to the long pBR322 sequence; Pyk3 having the promoter region proximal to the short pBR322 sequence and distal from the GAL4 sequence; while Pyk4 has the promoter region proximal to the GAL4 region, which in turn is proximal to the long pBR322 sequence; Pyk5 has the promoter region proximal to the GAL4 region which is proximal to the short pBR322 sequence; while Pyk6 has a promoter region distal from the GAL4 region and proximal to the long pBR322 sequence.

These plasmids described above were transformed into S. carlsbergensis strain 2150-2-3 (available from Lee Hartwell, University of Washington) under conventional conditions (Hinnen et al., Proc. Natl. Acad. Sci. USA (1978) 75:1929-1933). Cultures of 50-150ml were grown to mid or late log phase in rich media (YEP) under neutral conditions (3% glycerol, 2% lactic acid), and then inducing conditions (+2% galactose), or

repressing conditions (+2% glucose) for the final 1-2
generations. After lysis with glass beads and clarifi-
cation of the supernatants by centrifugation, HBsAg
expression was determined as described above. The re-
5 sults for the 12 plasmids are set forth in the following
Table 1.

Table 1: Expression of HBsAg from Gal Regulated Hybrid Promoters

	Construction	YEP + Glycerol / µg sAg/mg protein	YEP + Galactose µg sAg/mg protein	YEP + Glucose µg sAg/mg protein	Induction (Gal/glyc rol lactic acid)
5	GAP1	0.04	0.09	0.02	2.0
	GAP2	1.65	0.8	1.4	0.5
	GAP3	0.25	0.30	---	1.2
	GAP4	0.10	0.75	---	7.5
	GAP5	0.25	2.1	---	8.4
	GAP6	1.55	1.4	1.0	0.9
10	PYK1	0.10	0.30	0.14	3.0
	PYK2	1.65	1.4	1.1	0.85
	PYK3	0.10	0.15	---	1.5
	PYK4	0.10	1.0	0.05	10.0
	PYK5	0.03	1.4	0.02	47.0
	PYK6	1.7	1.8	0.9	0.9

Construction of pPGAP.

A yeast expression vector was prepared called pPGAP having a polyrestriction site linker between the GAPDH terminator and short promoter region. Plasmid plot5pGAP128 was digested with BamHI and TaqI to yield an approximately 390bp BamHI-TaqI fragment having the -400 to -26bp of the GAPDH promoter. The BamHI-TaqI fragment was ligated to a synthetic fragment having the following sequence:

10 CGA₂TA₃(CA)₃TA₃CA₃CACCATG₃A₂T₂CGT₂AG₂
 T₂AT₃(GT)₃AT₃GT₃GTGGTAC₃T₂A₂GCA₂TC₂AGCT

to provide a BamHI-SalI fragment, which was digested with BamHI and SalI and used to replace the BamHI-SalI fragment of BamHI-SalI digested pBR322 treated with alkaline phosphatase. After ligation, the plasmid pGAPNRS was obtained which was digested with BamHI and SalI to provide a 400bp BamHI-SalI fragment which was gel isolated. This fragment was ligated to an about 900bp SalI-BamHI fragment containing the GAPDH terminator region and a short segment of 3' coding region and the resulting 1.4kb BamHI-BamHI fragment digested with BamHI. The SalI-BamHI GAPDH terminator fragment was obtained by SalI and BamHI digestion of pGAP2, a plasmid prepared by insertion of an about 3.3kb BamHI fragment containing the GAPDH gene GAP49 (Holland and Holland, supra) into the BamHI site of pBR322. Plasmids pGAP2 and pGAP1 were obtained as follows: A yeast gene library was prepared by inserting fragments obtained after partial digestion of total yeast DNA with restriction endonuclease Sau3A in lambda-phage Charon 28 (Blattner et al., Science (1977) 196:161-169). The phage library was screened with DNA complementary to the yeast GAPDH mRNA and the yeast GAPDH gene from one of these clones was subcloned as either an about 3.3kb BamHI fragment in the BamHI site of pBR322 (pGAP-2) or as an about

2.1kb HindIII fragment in the HindIII site of pBR322 (pGAP-1).

pBR322 was digested with EcoRI and SalI, the termini blunt-ended and ligated to BamHI linkers, followed by BamHI digestion and the BamHI-BamHI 3.8kb fragment gel isolated, recircularized by self-ligation, cloned and designated pBRAR1-Sal. The 1.4kb BamHI-BamHI fragment was inserted into the BamHI-digested, alkaline phosphatase treated pBRAR1-Sal vector to provide the plasmid pPGAP of about 5.3kb with the orientation in the opposite direction of the amp^r.

The plasmid phSOD was prepared as follows:

Molecular cloning of hSOD cDNA.

Total RNA was prepared from an adult human liver by the guanidinium thiocyanate/lithium chloride method (Cathala *et al.*, DNA (1983) 2:329-335). polyA RNA was used to synthesize double-stranded cDNA (Maniatis *et al.*, Molecular Cloning, 213-242, Cold Spring Harbor, 1982) and this was passed over a Sepharose CL4B column to enrich for cDNAs of greater than 350bp (Fiddes and Goodman, Nature (1979) 281:351-356). The cDNA was inserted at the PstI site of plot4, a pBR322 derivative having the following sequence replacing the PstI-EcoRI site.

```

25      PstI      HinfI      AluI
      1      GGTGAATCCGTAATCATGGTCATAGCTGTTTCCTGTGTGAAATTGTTATCCGCTC
          ACGTCCACTTAGGCATTAGTACCAAGTATCGACAAAGGACACACTTTAACAATAGGCGAG
          HphI

      30      60      HindIIIAluI
          ACAATTCCACACATTATACGAGCCGATGATTAATTGTCAACAGCTCATTTCAGAATATTT
          TGTTAAGGTGTGTAATATGCTCGGCTACTAATTAACAGTTGTGAGTAAAGTCTTATAAA

          EcoRI
120      GCCAGAACCGTTATGATGCGG
          CGGTCTTGGCAATACTACGCCTTAA

```

The cDNA insert employed the lig -dG:dC tailing method (Maniatis et al., supra). E. coli strain D1210 was transformed with this mixture and transformants selected on L-agar containing 10µg/ml tetracycline

5 (Kushner, S.R. (1978) In: Genetic Engineering, eds. Boyer, H.B. and Nicosia, S., (Elsevier/North Holland, Amsterdam) p. 17). Plasmid DNA constituting a liver cDNA library was prepared (Maniatis et al., Molecular Cloning, pp. 86-94, Cold Spring Harbor 1982) directly

10 from approximately 62,000 recombinant colonies plated at a density of approximately 3,000 colonies per 9cm diameter Petri dish.

Isolation of r-hSOD clones.

Strain D1210 was retransformed with the liver

15 cDNA library and about 40,000 clones were grown on nine 14cm diameter Petri dishes. After transfer of the colonies to nitrocellulose paper and chloramphenicol amplification of plasmid DNA, the cells were lysed and the filters prepared for hybridization (Ish-Horowicz and

20 Burke, Nucleic Acids Research (1981) 9:2989-2998). Oligonucleotide probes were employed for screening by hybridization, with the probes consisting of enzymatically-radiolabeled, chemically-synthesized DNA molecules complementary to the mRNA encoding amino acid residues

25 19 to 24 of the protein (Jabusch et al., supra.; Barra et al., supra.); the mixture had the following sequences:

3' TTA AAA CTT GTT TTT CT 5'
 G G C C C

where all of the indicated possibilities for encoding

30 the peptide sequence were prepared (32-fold degenerate).

The probes were labeled with ³²P to a specific activity of 1-3x10⁸ cpm/µg and Millipore (0.45µm) filter used before use. Filters were prehybridized for 6hrs at 30°C in 4x SSC, 2x Denhardt's solution, 40mM sodium

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part of the variable linker segment (s discussi n relating to Fig. 3).

Construction of plot5 derivativ s containing r-hSOD.

The synthetic DNA molecules F(26), C(16),
5 B(31), D(11), E(13) and 4(24) shown in Fig. 3, were synthesized by the phosphoramidite method.

The single strand 4(24) was prepared by using all four bases, at each site where X is indicated. Furthermore, silica was withdrawn from the synthesis of
10 the 24mer, such that single-stranded 21mers, 22mers, and 23mers are obtained in addition to the 24mers. After removal from the silica support, the four mixtures are combined in appropriate proportions to provide f r equimolar amounts of each of the possible single strands.
15 This mixture was treated as a single product in the subsequent steps.

Molecules F(26), C(16), B(31) and D(11) were mixed together in equimolar amounts and 10µg phosphorylated using T4 polynucleotide kinase. After phenol-ether
20 extraction, the additional non-phosphorylated synthetic DNA molecules 4(24) and E(13) were added, such that all fragments were equimolar. The equimolar mixture contained 13µg of DNA in 133µl of 0.3x kinase buffer.

After annealing by cooling at a uniform rate
25 from 70°C to 20°C over 60min, the single strands were ligated together with T4 ligase in 200µl ligation mix at 14°C for 4hrs, phenol-chloroform extracted, ethanol precipitated and the 5'-ends of 4(24) and E(13) phosph -
30 al., supra). Preparative polyacrylamide gel electr - phoresis was used to isolate the completely ligated 53bp material having 5'- and 3'-overhangs.

The above purified fragment mixture was th n ligated to the 460bp TagI-PstI segment of the hSOD cDNA
35 as shown in Fig. 3. This segment was itself c nstructed by isolating the 454bp TagI-AluI hSOD fragment, making

it flush-ended using Klenow and inserting it into pl t5 between its EcoRI and SalI sites which had been similarly made flush-ended. After preparation of plasmid DNA from this recombinant, the 460bp TaqI-PstI hSOD fragment was isolated by preparative polyacrylamide gel electrophoresis. After extraction and precipitation, the 515bp fragment resulting from the joining of the synthetic fragment to the 460bp TaqI-PstI hSOD fragment was blunt-ended (525-528bp) and then digested with SalI and the resulting 519-522bp hSOD fragment isolated by polyacrylamide gel electrophoresis. This fragment was then inserted into plot5 which had been digested with PvuII and SalI and then treated with alkaline phosphatase. The resulting plasmids were used to transform strain D1210. Recombinants obtained after transformation of strain D1210 were selected on L-agar containing 100µg/ml ampicillin to give a set of clones, which were screened for an NcoI site. One was selected and designated phSOD.

20 Construction of a yeast vector for SOD expression.

The plasmid phSOD was digested with NcoI and SalI and a 550bp fragment obtained, which included 1 nucleotide untranslated at the 5'-terminus and the entire coding region for hSOD. pPGAP was digested with NcoI and SalI followed by treatment with alkaline phosphatase and the SalI-NcoI fragment substituted for the NcoI-SalI fragment in pPGAP to provide pPGAPSOD. BamHI digestion of pPGAPSOD resulted in a 2kb fragment which was gel isolated and inserted into the BamHI site of pCl/1 and pCl/1 GAL4/370. These plasmids were transformed into yeast strain 2150-2-3 as described previously, with the results of expression set forth in the following Table 2.

Table 2: Expression of Human SOD in Yeast Strain 2150

5	Plasmid	Carbon Source	SOD ² µg/mg protein
	pC1/1	g, L ¹	0
	pC1/1GAPSOD	g, L	148
10	pC1/1GALGAPSOD	g, L gal	0.4 68

15

¹ All cultures grown in Minus Leucine media with 2% lactic acid, 3% glycerol with or without 2% galactose to late log or early stationary phase.

² Determined by RIA.

20

hSOD levels were measured using a standard radioimmunoassay with iodinated authentic hSOD as standard. Constitutive synthesis from the GAP promoter leads to very high levels of hSOD production, of the order of 10-30% of the total cell protein. The induction with galactose works almost as well, yielding about 7% of the cell protein as hSOD.

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Construction of pYAS11

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This yeast expression plasmid contains the hSOD gene fused to the amino terminus of the human pro-insulin gene, with a methionine codon at the junction between both genes. The fusion gene is under control of the hybrid inducible ADH2-GAP (yeast alcohol dehydrogenase 2) promoter and the GAP terminator. An about 3kbp BamHI expression cassette was constructed.

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The ADH2 portion of the promoter was constructed by cutting a plasmid containing the wild type ADH2 gene (plasmid pADR2, see Beier and Young, Natur (1982) 300:724-728), a pBR322 derivative, which contains the yeast centromere gene from chromosome 3 (CEN3) (U.S. Patent No. 4,464,472), the Trp1 gene and the yeast ADHI1

gene with the restriction enzyme EcoR5, which cuts at a position +66 relative to the ATG start codon, as well as in two other sites in pADR2, outside of the ADH2 region in the CEN3 and Trp1 gene. The resulting mixture of a vector fragment and two smaller fragments was resected with Bal31 exonuclease to remove about 300bp. Synthetic XhoI linkers were ligated onto the Bal31 treated DNA. The resulting DNA linker vector fragment was separated from the linkers by column chromatography, cut with the restriction enzyme XhoI, religated and used to transform E. coli to ampicillin resistance. The positions of the XhoI linker additions were determined by DNA sequencing. One plasmid which contained an XhoI linker located within the 5' non-transcribed region of the ADH2 gene (position -232 from ATG) was cut with the restriction enzyme XhoI, treated with nuclease S1, and subsequently treated with the restriction enzyme EcoRI to create a linear vector molecule having one blunt end at the site of the XhoI linker and an EcoRI end at the pBR322 proximal terminus.

The GAP portion of the promoter was constructed by cutting plasmid pPGAP (supra) with the enzymes BamHI and EcoRI, followed by the isolation of the 0.4Kbp DNA fragment. The purified fragment was cut with the enzyme AluI to create a blunt end near the BamHI site.

Plasmid pJS014 was constructed by the ligation of the AluI-EcoRI GAP promoter fragment to the ADH2 fragment present on the linear vector described above.

Plasmid pJS104 was digested with BamHI (which cuts upstream of the ADH2 region) and with NcoI (which cuts downstream of the GAP region). The about 1.3Kbp fragment containing the ADH2-GAP promoter was gel purified and ligated to an about 1.7Kbp fragment containing the hSOD-proinsulin fusion DNA sequences and GAP terminator present in pYS11. This 3Kbp expression cassette was cloned into BamHI digested and phosphatase treated pCl1 to yield pYAS11.

Plasmid pYS11 was constructed as follows. Three fragments were employed which involve a 454bp NcoI-Sau3A isolated from phSOD, where the fragment includes the entire coding sequence for human superoxide dismutase (hSOD) with the exception of the last three 3'-codons; a 51bp Sau3A-HindIII synthetic adapter, which codes for the last three codons of hSOD, methionine, and the first 14 codons of proinsulin; and a 231bp HindIII-SalI fragment, isolated from pINS5 (a pBR322-derived vector which contains a proinsulin coding sequence chemically synthesized according to the amino acid sequence reported by Bell *et al.*, *Nature* (1979) 282:525-527), which encodes proinsulin excepting the first 14 amino acids. These fragments were ligated together and introduced into the plasmid pPGAP, which had been previously digested with NcoI and SalI and alkaline phosphatase treated. The resulting plasmid pS11 was digested with BamHI to provide an expression cassette which was cloned into plasmid pC1/1 to yield pYS11.

Cloning of alpha-1-antitrypsin.

A cDNA library was made from 10µg of polyA⁺ RNA isolated from a part of a human liver. This library was prepared by oligo-dT priming of the first cDNA strand and self-priming of the second cDNA strand. The ds cDNA was size fractionated on a Sepharose CL4B column and those molecules greater than 300bp isolated. This fraction was treated with nuclease S1 and tailed with dCTP, using terminal transferase. The tailed cDNA was annealed to pBR322 which had been digested with PstI and tailed with dGTP. Transformation of *E. coli* HB101 yielded 60,000 colonies, where greater than 90% of the clones were recombinant.

Two synthetic oligonucleotide probes were used to isolate the alpha-1-antitrypsin (α_1 -AT) cDNA, the first probe corresponding to amino acid residues

344-350 near the C-terminal of the protein was used to probe 5,000 colonies and the second probe, corresponding to amino acid residues -23 to -17 (+1 being the first nucleotide of the first codon of the mature α_1 -AT) of the signal peptide, was used to probe 25,000 colonies. The probe sequences were taken from the partial nucleotide sequence described by Kurachi *et al.*, Proc. Natl. Acad. Sci. USA (1981) 78:6826; Leicht *et al.*, Nature (1982) 297:655). Approximately 3% of the colonies hybridized to the C-terminal probe and four hybridized to the N-terminal probe. The four N-terminal clones and 12 C-terminal clones were isolated and subjected to restriction analysis. From these, three overlapping clones which cover the entire cDNA were subjected to further study and were used to construct the full length cDNA clone.

The entire sequence of a composite full length cDNA derived from the three plasmids is as follows:

-10

Val Ser Trp Gly Ile Leu Leu Leu Ala Gly Leu Cys Cys Leu Val Pro Val Ser Leu
GTC TCG TCG GGC ATC CTC CTG CTG GCA GGC CTG TGC TGC CTG GTC CCT GTC TCC CTG

1 Glu Asp Pr Gln Gly Asp Ala Ala Gln Lys Tyr Asp Thr Ser His His Asp Gln Asp
1 GAG GAT CCC CAG GCA GAT GCT GCC CAG AAG ACA GAT ACA TCC CAC CAT GAT CAG GAT
--- -- BamHI

21 Pro Thr Phe Asn Lys Ile Thr Pro Asn Leu Ala Glu Phe Ala Phe Ser L u Tyr Arg
61 CCA ACC TTC AAC AAG ATC ACC CCC AAC CTG GCT GAG TTC GCC TTC AGC CTA TAC CGC

41 Leu Ala His Gln Ser Asn Ser Thr Asn Ile Phe Phe Ser Pro Val Ser Ile Ala Thr
121 CTG GCA CAC CAG TCC AAC AGC ACC AAT ATC TTC TTC TCC CCA GTG AGC ATC GCT ACA

61 Phe Ala Met Leu Ser Leu Gly Thr Lys Ala Asp Thr His Asp Glu Ile Leu Glu Gly
181 TTT GCA ATG CTC TCC CTG GGC ACC AAG GCT GAC ACT CAC GAT GAA ATC CTG GAG GGC

81 Asn Phe Asn Leu Thr Glu Ile Pro Glu Ala Gln Ile His Glu Gly Phe Gln Glu Leu
241 AAT TTC AAC CTC ACC GAG ATT CCG GAG GCT CAG ATC CAT GAA GGC TTC CAG GAA CTC

Arg(a,c)

101 His Thr Leu Asn Gln Pro Asp Ser Gln Leu Gln Leu Thr Thr Asp Gly(c)
301 CAT ACC CTC AAC CAG CCA GAC AGC CAG CTC CAG CTG ACC ACC GGC AAT GGC CTG TTC

121 Ser Glu Gly Leu Lys Leu Val Asp Lys Phe Leu Glu Asp Val Lys Lys Leu Tyr His
361 ACC GAG GGC CTG AAG CTA GTG GAT AAG TTT TTG GAG GAT GTT AAA AAG TTG TAC CAC

141 Glu Ala Phe Thr Val Asn Phe Gly Asp Thr Glu Glu Ala Lys Lys Gln Ile Asn Asp
421 GAA CCC TTC ACT GTC AAC TTC GGC GAC ACC GAA CAG GCC AAG AAA CAG ATC AAC GAT

161 Val Glu Lys Gly Thr Gln Gly Lys Ile Val Asp Leu Val Lys Glu Leu Asp Arg Asp
481 GTG CAG AAG GGT ACT CAA GGG AAA ATT GTG GAT TTG GTC AAG GAG CTT GAC AGA GAC

181 Val Phe Ala Leu Val Asn Tyr Ile Phe Phe Lys Gly Lys Trp Glu Arg Pro Phe Glu
541 GTT TTT GCT CTG GTG AAT TAC ATC TTC TTT AAA GGC AAA TCG GAG AGA CCC TTT GAA

Ala(b)

201 Lys Asp Thr Glu Glu Glu Asp Phe His Val Asp Gln Val Thr Thr Val Lys Val Pro
601 AAG GAC ACC CAG GAA GAG GAC TTC CAC GTG GAC CAG GTG ACC ACC GTG AAG GTG CCT
--- -- BstEI

221 Met Lys Arg Leu Gly Met Phe Asn Ile Gln His Cys Lys Lys Leu Ser Ser Trp Val
661 ATG AAG CGT TTA GGC ATG TTT AAC ATC CAG CAC TGT AAG AAG CTG TCC AGC TGG GTG

Asn(c)

241 Leu Met Lys Tyr Leu Gly Asn Ala Thr Ala Ile Phe Phe Leu Pro Asp Glu Gly Lys
721 CTG ATG AAA TAC CTG GGC AAT GCC ACC GCC ATC TTC TTC CTG CCT GAT GAG GGG AAA

261 Gln His Leu Glu Asn Glu Leu Thr His Asp Ile Ile Thr Lys Phe Leu Glu Asn Glu
781 CAG CAC CTG GAA AAT GAA CTC ACC CAC GAT ATC ACC AAG TTC CTG GAA AAT GAA
--- -- EcoRV

281 Arg Arg Ser Ala Ser Leu His Leu Pro Lys Leu Ser Ile Thr Gly Thr Tyr Asp Leu
841 AGA AGG TCT GCC ACC TTA CAT TTA CCC AAA CTG TCC ATT ACT GGA ACC TAT GAT CTG

Val(a,c)

301 Ser Ile Leu Gly Gln Leu Gly Ile Thr Lys Val Phe Ser Asn Gly Ala Asp Leu Ser
901 AGC ATC CTG GGT CAA CTG GGC ATC ACT AAG GTC TTC AGC AAT GGC GCT GAC CTC TCC

321 Val Thr Glu Glu Ala Pro Leu Lys Leu Ser Lys Ala Val His Lys Ala Val Leu Thr
961 GTC ACA GAG GAG GCA CCC CTG AAG CTC TCC AAG GCC GTG CAT AAC GCT GTG CTG ACC

341 Asp Glu Lys Gly Thr Glu Ala Ala Gly Ala Met Phe Leu Glu Ala Ile Pro Met Ser
1021 CAC GAG AAA GGC ACT GAA GCT GCT GGC GCC ATG TTT TTA GAG GCC ATA CCC ATG TCT

361 Pro Pro Glu Val Lys Phe Asn Lys Pro Phe Val Phe Leu Met Ile Glu Gln Asn Thr
1081 CCC CCC GAG GTC AAG TTC AAC AAA CCC TTT GTC TTC TTA ATG ATT GAA CAA AAT ACC
--- -- Aval

381 Ser Pro Leu Ph Met Gly Lys Val Val Asn Pro Thr Gln Lys OC
1141 TCT CCC CTC TTC ATG GGA AAA GTG GTG AAT CCC ACC CAA AAA TAA CTGCCTCTCGCTCCT
--- -- HinfI

AAT CCC ACC CAA AAA TAG
GGG TGG GTT TTT ATC AGCT
----- SalI

1201 CCTCCCCCTCCATCCCTGCCCCCTCCCTGGATGACATTAAAGAAGGGTTGAGCTGGAAAAAAAAAAAAAAAAAAAA

LEGEND

5 Nucleotide and predicted amino acid sequences
 of α_1 -AT cDNA. The reactive center motif at
 positions 358-359 is boxed. Subscripts to
 10 amino acids in parentheses identify differences
 between the subject protein sequence and those
 derived from (a) protein sequencing (Carrell
et al., 1982), (b) the cDNA of Woo et al.,
 [see Carrell et al., 1982]], and (c) the cDNA
 of Bollen et al., 1983. The synthetic DNA
 molecules used in the construction of the
BamHI to SalI fragment encoding the mature
 protein are shown as are the cDNA restriction
 sites used in this construction.

15 The above sequence was determined using the
 dideoxy sequencing method of Sanger et al., Proc. Natl.
Acad. Sci. USA (1977) 74:5463, in the M13 vectors of
 Messing et al., Nucleic Acids Res. (1981) 9:309. The
 differences at the nucleotide and amino acid level from
 20 the published cDNA sequences are shown.

Construction of the full length clone for
 expression of yeast began with three fragments isolated
 from cDNA clones: 1) a 630bp BamHI-BstEII fragment; 2)
 a 450bp BstEII-AvaI fragment; and 3) an 85bp AvaI-HinfI
 25 fragment. A synthetic adapter was employed having the
 following sequence:

$$\begin{array}{c} A_2TC_3AC_3A_5TAG \\ G_3TG_3T_5ATCAGCT \end{array}$$

30 Approximately two pmoles of fragments 1 and 2 were li-
 gated together and after removal of the ligase, digested
 with BamHI and AvaI. Fragment 3 and the synthetic adap-
 ter were ligated and digested with AvaI and SalI and
 the two resulting fragment mixtures were ligated followed
 by digestion with BamHI and SalI. Fragments migrating
 35 upon electrophoresis in the region of about 1000-1400bp
 were isolated and cloned by substitution into BamHI and

SalI digest d and alkaline phosphatase treated pBR322.
The resulting pl smid is r ferr d to as pATi.

Plasmid pPGAP was digested with Nc I, f llowed
by blunt-ending, followed by SalI digesti n and treat-
5 ment with alkaline phosphatase. The NcoI-SalI fragment
was substituted with an approximately 1250bp blunt-ended
(BamHI)-SalI fragment obtained from plasmid pATi, by
BamHI digestion, blunt ending, and SalI digestion.
This was inserted into the pPGAP vector to produce the
10 plasmid pGAPATi, a 6.6kb plasmid, which was digested
with NcoI and BamHI and a 2.3kb NcoI-BamHI fragment
obtained having the α_1 -AT gene and the GAPDH terminator
and approximately 400bp BamHI-NcoI fragment obtained
having the GAPDH promoter. These fragments were ligated
15 together and inserted into the BamHI site of pCl/1.
The plasmids pCl/1GAPATi8 and pCl/1GAPATi9 were obtained
with the orientation of expression clockwise in the
former and counterclockwise in the latter, with amp^r
being in the counterclockwise direction. These plasmids
20 were transformed in S. cerevisiae AB103 (A.T.C.C. No.
20658, deposited January 5, 1983) by standard methods,
selecting for leucine prototrophy and grown as described
above. Yeast extracts were prepared by lysis with glass
beads and the α_1 -AT activity determined by inhibition
25 of human leukocyte elastase.

Assays contained in 1ml:0.1-0.2 human leukocyte
elastase (HLE); 0.1mM MeO-Suc-Ala-Ala-Pro-Val-p-nitr -
anilide (Beatty et al., J. Biol. Chem. (1980) 255:3931);
50mM Tris, pH 8, 0.5M NaCl, and the indicated amounts
30 of yeast extract or human α_1 -AT. Assays were initiated
by the addition of elastase, incubated at 28°C for 15min,
terminated by the addition of 100 μ l of 8N acetic acid
and the absorbance at 410nm determined. Typical results
are shown in the following Table 3.

Table 3

Plasmid	Strain	Amt.Extract (μ l)	Amt.HLE (μ g)	Amt.Protein (μ g)	% Elastase Activity	% α_1 -AT*
pCl/1GAPAT18	AB103	5.0	0.1	50.0	40	0.17
		10.0	0.1	100.0	26	0.11
pCl/1GAPAT19	AB103	0.25	0.1	2.3	89	0.7
		1.0	0.1	9.1	26	1.2
pCl/1GAPAT19	AB110	0.2	0.2	2.9	39	6.1
		0.4	0.2	5.8	14	4.3

* Calculation based upon the Mol. wt. of HLE (29kD), the amount of protein added and the degree of inhibition.

Th above data demonstrate that plasmids having the orientation of the expression cassette in the counter-clockwise orientation, the promoter proximal to the long sequence of pBR322, make 10-20 times more α_1 -AT than the same cassette in the other orientation.

Yeast strain AB110.

Yeast strain 2150-2-3 was crossed with a yeast strain AB103 transformant containing pC1/1GAPATi9. The diploids were sporulated and the tetrads dissected. Strains were maintained on leucine selective plates in order to ensure maintenance of the plasmid, since the parents are auxotrophs. A series of colonies were screened for their genotype with respect to a number of markers. The most vigorous strains were selected and cultures grown on leucine selective media. The best strain was designated AB110 (pC1/1GAPATi9), gave 6-7.5% of the total cell protein as α_1 -AT as shown in the above Table 3. The strain AB110 has the following genotype : Mat α , ura3-52, leu2-04 or both leu2-3 and leu2-112, pep4-3, his4-580 (cir o).

Phosphate induction.

Plasmid pPGT80 was digested with BamHI, the ends blunt-ended, followed by digestion with XbaI and the 500bp fragment containing the GAPDH promoter and 5'-end of the sAg gene isolated.

The PHO5 gene was isolated from a yeast genomic library employing an oligonucleotide probe 5'-GGCACTCACACGTGGGACTAG-3' derived from the published partial sequence (Meyhack et al., The EMBO Journal (1982) 1:675-680). A subfragment of this clone containing 550bp of the 5'-untranslated region and approximately 80bp of coding sequence was subcloned as a BamHI-SalI substitution in pBR322 to provide pPHO5. This fragment has the sequence 5'-ATGTTTAAA-3', encoding the first three amino acids, the second and third codons

specifying an AhaIII site. The plasmid pHBS6 was digested with NcoI, blunt-ended, followed by digestion with BamHI and treatment with alkaline phosphatase. The PHO5 promoter region was obtained by digesting the pPHO5 plasmid with AhaIII, resecting the resulting fragment with Bal31 for a short time, followed by digestion with BamHI and isolation of a 500-550bp BamHI blunt-ended fragment. This fragment was employed for substitution of the NcoI-BamHI fragment from pHBS6 and was screened for regeneration of the NcoI restriction site to provide plasmid pHBS6PHO5/1.

Plasmid pHBS6PHO5/1 was digested with BstEII which cleaves at position -175 in the PHO5 promoter. This molecule was blunt-ended, digested with SalI and the 650bp fragment having the 5'-portion of the promoter domain, containing 275bp of pBR322 and 375bp of the PHO5 promoter region isolated. This fragment was ligated with the blunt-ended (BamHI)-XbaI fragment obtained from digestion of pPGT80 with BamHI, blunt ending, followed by XbaI digestion. After digesting the ligated fragment with SalI and XbaI, the resulting fragment was then substituted into pPGT16-3 which had been digested with SalI and XbaI and treated with alkaline phosphatase. The resulting plasmid pPHO5PGT80 had a cassette comprising the PHO5 regulatory region, the GAPDH promoter, the sAg gene and the ADH terminator. This cassette was excised from the plasmid by BamHI digestion, whereby a 1.8kb BamHI-BamHI fragment was gel isolated and ligated into the BamHI site of BamHI digested and alkaline phosphatase treated pCl/1 to provide plasmids PHO5GAP1 and PHO5GAP2 where the PHO5 was distal and proximal to the long pBR322 sequence, respectively.

The two plasmids were transformed into yeast strain 2150-2-3 as described above and grown in rich media as described above for 8 to 10 generations in either high (7mM) or low (0.2mM) phosphate. Samples were harvested in late log phase and HBsAg determined

as described previously. The results are shown below in Table 4.

Table 4: Regulation of HBsAg Production in Yeast using a Hybrid PHO5/GAPDH Promoter.

Construction	High Phosphate (7mM) (sAg µg/mg protein)	Low Phosphate (0.2mM)	Induction low / high
PHO5GAP-1	0.08	0.95	12.0
PHO5GAP-2	0.27	0.40	1.5

From the above results, it is evident that effective regulation with phosphate is obtained, with one orientation being superior to the other.

It is evident from the above results, that highly efficient expression can be obtained, either constitutive or regulated, by providing for truncated promoter regions of yeast glycolytic enzyme gene promoters, employing the 3' domain proximal to the coding region of the gene in conjunction with a 5'-portion or second domain of the promoter region of a yeast gene subject to inducible regulation by a nutrient, e.g., carbon source or phosphate, temperature, or other externally controllable source or condition. Alternatively, the second domain may be replaced by prokaryotic sequences of at least about 1kb or greater, which provide for constitutive enhancement in the absence of the second domain of the promoter region. Thus, a wide variety of genes exogenous to yeast may be expressed in high yield in high percentages of the total protein of the yeast host.

Alth ough the f regoing invention has b en
d scribed in some detail by way of illustration and
exampl for purp ses of clarity of understanding, it
will be obvious that certain chang s and modifications
5 may be practiced within the scope of the appended claims.

WHAT IS CLAIMED IS:

1. In a method for obtaining expression of a foreign structural gene in a yeast host employing hybrid constructions involving transcriptional and translational regulatory signals recognized by said host joined to a foreign gene for transcriptional and translational regulation of expression in said host, wherein said construction is introduced into said host and said foreign gene is expressed to produce the encoded polypeptide,
 - the improvement which comprises:
 - employing for transcriptional regulation, a DNA sequence having first and second domains,
 - said first domain being proximal to said structural gene, being of less than about 600bp and having a sequence which includes the RNA polymerase binding site and transcription initiation site of a yeast glycolytic enzyme gene; and
 - said second domain distal to said structural gene providing for enhanced efficiency of transcription and being the regulatory region of a regulated yeast gene other than the natural gene of said first domain or a prokaryotic sequence of at least about 500bp.
2. A method according to Claim 1, wherein said yeast glycolytic enzyme gene of said first domain is glyceraldehyde-3-phosphate dehydrogenase.
3. A method according to Claim 2, wherein said regulatory region is the GAL1 or GAL10 regulatory region.
4. A method according to Claim 2, wherein said regulatory region is the PHO5 regulatory region.

5. A method according to Claim 1, wherein said yeast glycolytic enzyme of said first domain is pyruvate kinase.

6. A method according to any of Claims 1, 2, 3, 4, or 5, wherein said polypeptide is Hepatitis B surface antigen, superoxide dismutase, or α_1 -antitrypsin.

7. In a method for obtaining expression of a foreign structural gene in a yeast host employing a hybrid construction involving transcriptional and translational regulatory signals recognized by said host joined to a foreign gene for transcriptional and translational regulation of expression in said host, wherein said construction is introduced into said host and said foreign gene is expressed to produce the encoded polypeptide, the improvement which comprises employing for transcriptional regulation, a DNA sequence having first and second domains,

said first domain being proximal to said structural gene, being of about 200-500bp, being derived from the 5'-untranslated region of the yeast glyceraldehyde-3-phosphate dehydrogenase or pyruvate kinase gene and extending downstream to at least nucleotide -10; and

said second domain distal to said structural gene, being a prokaryotic sequence of at least about 1000bp, wherein said yeast cells containing said construct are grown to high density; and isolating said polypeptide.

8. In a method for obtaining expression of a foreign structural gene in a yeast host employing hybrid constructions involving transcriptional and translational regulatory signals recognized by said host joined to a foreign gene for transcriptional and translational regulation of expression in said host, wherein

said constructi n is introduced into said host and said foreign g ne is expressed to produc the encoded polypeptide, th improvement which c mprises employing for transcriptional regulation, a DNA sequence having first
5 and second domains,

said first domain being proximal to said structūral gene, being of about 200-500bp and including the RNA polymerase binding site and transcription initiation site of the yeast glyceraldehyde-3-phosphate dehydrogenase or pyruvate kinase gene; and
10

said second domain distal to said structural gene having a sequence providing for regulation of th acid phosphatase PHO5 gene or the GAL1 or GAL10 gene, growing said host containing said construct
15 in a nutrient medium to high density of said host under non-inducing conditions;

changing the conditions to inducing conditions to induce expression of said structural gene; and isolating the polypeptide.

20 9. A DNA construct comprising in the downstream direction a yeast promoter region having first and second domains, an insertion sequence of from ab ut 4-100bp having at least one unique restriction site for the construct, and a termination region, wherein said
25 two domains are a distal domain of at least about 100bp of a DNA sequence providing for regulation of a yeast gene or a prokaryotic sequence of at least about 500bp and a proximal domain which includes the RNA polymerase binding site and the transcription initiation site f a
30 yeast glycolytic enzyme gene.

10. A DNA construct for expression of a structural gene foreign to yeast, comprising in the downstr am dir ction:

a promoter region comprising two domains and providing for transcriptional and translational signals for initiation in yeast;

5 a structural gene under regulatory control of said promoter region; and

a termination region providing for transcriptional and translational termination in yeast;

10 wherein the distal domain comprises a sequence of at least about 100bp including a sequence for regulation of a regulatable yeast gene or a prokaryotic sequence of at least about 500bp and the proximal domain includes an RNA polymerase binding site and transcription initiation site of a yeast glycolytic enzyme gene.

11. A construct according to Claim 10, wherein said gene is α_1 -antitrypsin, superoxidase dismutase or Hepatitis B surface antigen.

12. A modified yeast host containing as an extrachromosomal element a plasmid vector having a replication system, at least one marker, and a cassette
20 comprising in the downstream direction of transcription, a promoter region functional in yeast for transcriptional and translational initiation, a structural gene foreign to said yeast under the transcriptional and translational control of said promoter region, and a terminator region
25 functional in said yeast, wherein said promoter region has two domains, a distal domain distant from said gene of at least about 100bp comprising a sequence providing for regulation of a regulated yeast gene or a prokaryotic sequence of at least about 500bp, and a proximal
30 domain proximal to said gene, comprising the RNA polymerase binding site and transcription initiation site of a yeast glycolytic enzyme gene and of about 200 to 600bp and extending to at least the -20nt of said yeast glycolytic enzyme gene.

13. A yeast cell according to Claim 12, wherein the yeast glycolytic enzyme for said proximal domain is glyceraldehyde-3-phosphate dehydrogenase or pyruvate kinase.

5 14. A method for producing a foreign protein which comprises:

growing cells according to Claim 12, wherein said foreign protein is produced.

10 15. A method according to Claim 14, wherein said foreign protein is α_1 -antitrypsin, human superoxide dismutase or Hepatitis B surface antigen.

16. A method according to Claim 2, wherein said regulatory region is the ADHII regulatory region.

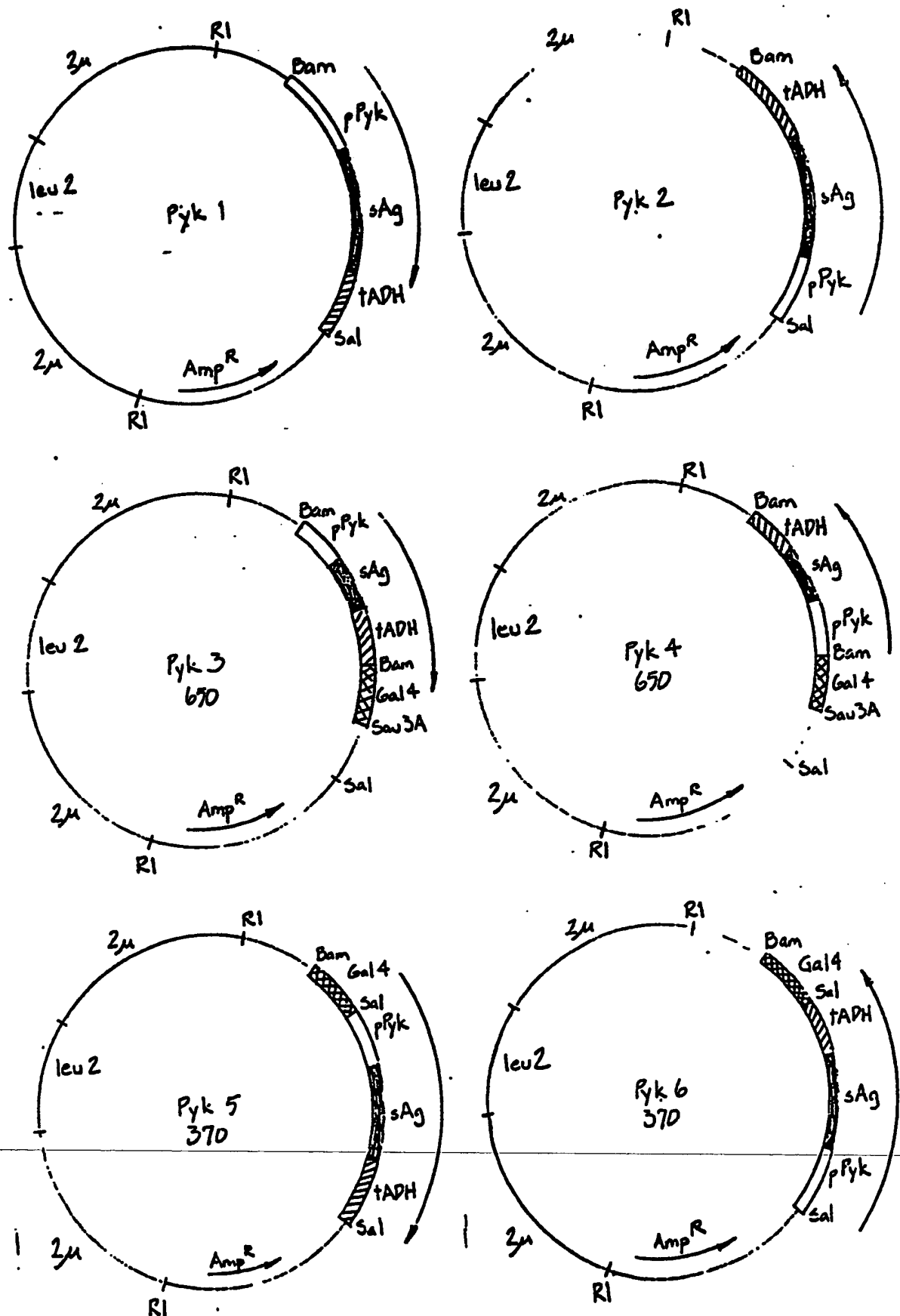


FIG. 2.

FIG.—3.

